

U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE FORM PTO-1390 (REV 12-28-99)		ATTORNEY'S DOCKET NUMBER
TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371		GJE-39
INTERNATIONAL APPLICATION NO. PCT/GB98/02628 ✓	INTERNATIONAL FILING DATE September 2, 1998 ✓	U.S. APPLICATION NO. (If known) see 37 CFR 1.5 09/486676 ✓
TITLE OF INVENTION <u>Conjugates That Contain the Homeodomain of Antennapedia</u>		
APPLICANT(S) FOR DO/EO/US <u>Andrea Crisanti</u>		
Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:		
<ol style="list-style-type: none"> <input checked="" type="checkbox"/> This is a FIRST submission of items concerning a filing under 35 U.S.C. 371. <input type="checkbox"/> This is a SECOND or SUBSEQUENT submission of items concerning a filing under 35 U.S.C. 371. <input checked="" type="checkbox"/> This express request to begin national examination procedures (35 U.S.C. 371(f)) at any time rather than delay examination until the expiration of the applicable time limit set in 35 U.S.C. 371(b) and PCT Articles 22 and 39(1). <input checked="" type="checkbox"/> A proper Demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date. <input checked="" type="checkbox"/> A copy of the International Application as filed (35 U.S.C. 371(c)(2)) <ol style="list-style-type: none"> a. <input type="checkbox"/> is transmitted herewith (required only if not transmitted by the International Bureau). b. <input checked="" type="checkbox"/> has been transmitted by the International Bureau. c. <input type="checkbox"/> is not required, as the application was filed in the United States Receiving Office (RO/US). <input type="checkbox"/> A translation of the International Application into English (35 U.S.C. 371(c)(2)). <input type="checkbox"/> Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3)) <ol style="list-style-type: none"> a. <input type="checkbox"/> are transmitted herewith (required only if not transmitted by the International Bureau). b. <input type="checkbox"/> have been transmitted by the International Bureau. c. <input type="checkbox"/> have not been made; however, the time limit for making such amendments has NOT expired. d. <input type="checkbox"/> have not been made and will not be made. <input type="checkbox"/> A translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)). <input checked="" type="checkbox"/> An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)). (unsigned) <input type="checkbox"/> A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)). 		
Items 11. to 16. below concern document(s) or information included:		
<ol style="list-style-type: none"> <input type="checkbox"/> An Information Disclosure Statement under 37 CFR 1.97 and 1.98. <input type="checkbox"/> An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included. <input checked="" type="checkbox"/> A FIRST preliminary amendment. <input type="checkbox"/> A SECOND or SUBSEQUENT preliminary amendment. <input type="checkbox"/> A substitute specification. <input type="checkbox"/> A change of power of attorney and/or address letter. <input type="checkbox"/> Other items or information: 		

U.S. APPLICATION NUMBER (37 CFR 1.47)		INTERNATIONAL APPLICATION NO. PCT/GB98/02628	ATTORNEY'S DOCKET NUMBER GJE-39																							
09/486676		CALCULATIONS PTO USE ONLY																								
<p>17. <input checked="" type="checkbox"/> The following fees are submitted:</p> <p>BASIC NATIONAL FEE (37 CFR 1.492 (a) (1) - (5)):</p> <p>Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO \$970.00</p> <p>International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO \$840.00</p> <p>International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.445(a)(2)) paid to USPTO \$690.00</p> <p>International preliminary examination fee paid to USPTO (37 CFR 1.482) but all claims did not satisfy provisions of PCT Article 33(1)-(4) \$670.00</p> <p>International preliminary examination fee paid to USPTO (37 CFR 1.482) and all claims satisfied provisions of PCT Article 33(1)-(4) \$96.00</p>																										
ENTER APPROPRIATE BASIC FEE AMOUNT = \$ 840.00																										
<p>Surcharge of \$130.00 for furnishing the oath or declaration later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(e)).</p> <table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th>CLAIMS</th> <th>NUMBER FILED</th> <th>NUMBER EXTRA</th> <th>RATE</th> </tr> </thead> <tbody> <tr> <td>Total claims</td> <td>22</td> <td>- 20 =</td> <td>2 X\$18.00</td> </tr> <tr> <td>Independent claims</td> <td>8</td> <td>- 3 =</td> <td>5 X\$78.00</td> </tr> <tr> <td colspan="3">MULTIPLE DEPENDENT CLAIM(S) (if applicable)</td> <td>+ \$260.00</td> </tr> <tr> <td colspan="4" style="text-align: right;">TOTAL OF ABOVE CALCULATIONS = \$1,266.00</td> </tr> </tbody> </table>				CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE	Total claims	22	- 20 =	2 X\$18.00	Independent claims	8	- 3 =	5 X\$78.00	MULTIPLE DEPENDENT CLAIM(S) (if applicable)			+ \$260.00	TOTAL OF ABOVE CALCULATIONS = \$1,266.00						
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<p>Reduction of 1/2 for filing by small entity, if applicable. A Small Entity Statement must also be filed (Note 37 CFR 1.9, 1.27, 1.28).</p> <table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th colspan="3">SUBTOTAL = \$1,266.00</th> </tr> </thead> <tbody> <tr> <td colspan="3">Processing fee of \$130.00 for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(f)).</td> </tr> <tr> <td colspan="3" style="text-align: right;">TOTAL NATIONAL FEE = \$1,266.00</td> </tr> <tr> <td colspan="3">Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property</td> </tr> <tr> <td colspan="3" style="text-align: right;">TOTAL FEES ENCLOSED = \$1,266.00</td> </tr> <tr> <td colspan="3"></td> <td style="text-align: right;">Amount to be refunded: \$</td> </tr> <tr> <td colspan="3"></td> <td style="text-align: right;">charged: \$</td> </tr> </tbody> </table>				SUBTOTAL = \$1,266.00			Processing fee of \$130.00 for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(f)).			TOTAL NATIONAL FEE = \$1,266.00			Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property			TOTAL FEES ENCLOSED = \$1,266.00						Amount to be refunded: \$				charged: \$
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<p>a. <input type="checkbox"/> A check in the amount of \$_____ to cover the above fees is enclosed.</p> <p>b. <input checked="" type="checkbox"/> Please charge my Deposit Account No. <u>19-0065</u> in the amount of \$<u>1,266.00</u> to cover the above fees. A duplicate copy of this sheet is enclosed.</p> <p>c. <input checked="" type="checkbox"/> The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. <u>19-0065</u>. A duplicate copy of this sheet is enclosed.</p>																										
<p>NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137(a) or (b)) must be filed and granted to restore the application to pending status.</p>																										
<p>SEND ALL CORRESPONDENCE TO:</p> <p>Doran R. Pace Saliwanchik, Lloyd & Saliwanchik A Professional Association 2421 N.W. 41st Street, Suite A-1 Gainesville, FL 32606</p>																										
 <p>SIGNATURE: Doran R. Pace NAME <u>38,261</u> REGISTRATION NUMBER</p>																										

March 1, 2000

PRELIMINARY AMENDMENT
Patent Application
Docket No. GJE-39

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant : Andrea Crisanti

Docket No. : GJE-39

For : Conjugates That Contain the Homeodomain of Antennapedia

Box PCT

Assistant Commissioner for Patents
Washington, D.C. 20231

PRELIMINARY AMENDMENT

Sir:

Please amend the above-identified patent application as follows:

In the Specification

After page 37: Please insert as new page 38 the attached Abstract of the Disclosure.

In the Claims

The following amendments are made with respect to the claims in the international application PCT/GB98/02628 attached as Annexes to the International Preliminary Examination Report (IPER). Therefore, please replace existing pages 36-38 of the international application with the amended claim sheets (replacement pages 1-2) of the annex attached to the IPER as new pages 36-37, and make the following amendments to the pending claims:

Claim 2, line 1: Delete "A" and insert --The--.

Claim 3, line 1: Delete "A" and insert --The--.

Claim 4, line 1: Delete "A" and insert --The--.

Claim 4, line 1: Delete "any preceding claim" and insert --claim 1--.

Claim 5, line 1: Delete "A" and insert --The--.

Claim 6, line 1: Delete "A" and insert --The--.

Claim 7, line 1: Delete "A" and insert --The--.

Claim 8, line 1: Delete "A" and insert --The--.

Claim 9, line 1: Delete "A" and insert --The--.

Claim 9, line 1: Delete "or claim 2".

Claim 10, line 1: Rewrite "Nucleic acid" as --A nucleic acid--.

Claim 10, lines 1-2: Delete "any preceding claim" and insert --claim 1--.

Claim 13, lines 1-2: Delete "any of claims 1 to 9" and insert --claim 1--.

Claim 14, line 1: Delete "A" and insert --The--.

Claim 16, line 2: Delete "any one of claims 1 to 9 or claim 15" and insert --claim 1--.

Claim 17, line 1: Delete "A" and insert --The--.

Claim 19, line 1: Delete "A" and insert --The--.

Claim 19, lines 1-2: Delete "any one of claims 1 to 9 or claim 15" and insert --claim 1--.

Please cancel claims 15 and 18, without prejudice.

Please add the following new claims 20-24:

1 20. The conjugate according to claim 7, wherein the second region is a histone
2 protein.

1 21. The conjugate according to claim 2, wherein the second region comprises an
2 NOI.

1 22. A conjugate prepared by a method comprising the steps:
2 (i) culturing a host cell according to claim 12 under conditions which provide for the
3 expression of the conjugate within the host cell; and
4 (ii) recovering the conjugate by affinity purification under non-denaturing conditions.

1 23. The conjugate prepared according to claim 22, wherein the conjugate comprises
2 an amino acid tail that binds to an immobilised substrate.

1 24. A method for the treatment or prevention of a condition selected from the group
2 consisting of cancer, genetic disease, bacterial infection and viral infection, said method
3 comprising administering an effective amount of a conjugate of claim 1 to a person or animal
4 in need of such treatment, wherein the second region of the conjugate comprises a moiety
5 useful in treating or preventing said condition.

The Commissioner is hereby authorized to charge any fees under 37 CFR 1.16 or 1.17 as required by this paper to Deposit Account 19-0065.

Respectfully submitted,



Doran R. Pace
Patent Attorney
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Gainesville, FL 32606

DRP/sl
Attachment: Abstract of the Disclosure

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CONJUGATES THAT CONTAIN THE HOMEODOMAIN OF ANTENNAPIEDIA

- 5 The present invention relates to a novel conjugate, particularly, but not exclusively, in the form of a fusion protein or protein/nucleic acid complex.

Gene therapy provides the potential to permanently cure selected genetic diseases. However, a major obstacle is the effective delivery of the gene or protein of interest 10 to the target site. A variety of viral and non-viral vectors have been developed to deliver genes or gene products to various cells, tissues and organs by *ex vivo* or *in vivo* strategies. Among viral-based vectors, retroviruses, adenoviruses, adeno-associated viruses and herpes viruses have been most extensively studied. Among 15 non-viral-based vectors, liposomes have been used to introduce plasmid DNA directly into animals. However, one of the main challenges of gene therapy remains the design of effective delivery systems.

The gene *antennapedia* (*Antp*) encodes a transcriptional factor that has been shown to control antero-posterior morphogenesis in *Drosophila* embryo. The protein sequence 20 of *antennapedia* is characterised by the presence of a 60 amino acids motif (homeodomain) that binds to specific DNA target elements. *Antennapedia* homologues have been found in nearly all multicellular organisms and show a very high degree of amino acid sequence identity. The human and *drosophila* 25 *antennapedia* proteins differ in the sequence of the homeodomain only for one conservative amino acid substitution.

It has been observed that *antennapedia* and its homeodomain are able to translocate across the cytoplasmic membrane of mammalian cells. The translocation does not depend on cell endocytosis and it has been reported that translocation occurs at both

4°C and 37°C. Homeodomain synthetic peptides made of D amino acids are also able to cross the cytoplasmic membrane. This finding would rule out the possibility that

Antp is translocated through a receptor mediated mechanism. This property has been exploited to vehiculate small viral sequences into the cytoplasm of cultured cells as

- 5 well as to elicit an MHC class I restricted cytotoxic immune response against the nucleoprotein of the influenza virus. However, to date the homeodomain of *Antp* has only been used to transport small synthetic peptides. Schutze-Redelmeier M-P et al (1996) Introduction of Exogenous Antigens into the MHC Class I Processing and Presentation Pathway by *Drosophila* Antennapedia Homeodomain Primes Cytotoxic

- 10 T Cells In Vivo. *The Journal of Immunology* 650-655 mentions that the homeodomain of the antennapedia molecule can be used to deliver up to 50 additional amino acids to the cytoplasm. Using 16 amino acids of the third helix they mention that fusion peptides containing up to 94 amino acids could be prepared, but this is not demonstrated. Further the thrust of the disclosure is only towards delivery 15 of synthetic peptides.

We have now found that the homeodomain of *Antp* can be used to translocate proteins, including non-synthetic proteins. One of the key advantages of the present invention is that it can be used to translocate functional and regulatory proteins. This

- 20 is surprising and very important, particularly for medical applications.

The ability to deliver the expression product of a gene of interest directly into a cell has wide applicability, particularly in the medical field. We have now also surprisingly found that the homeodomain is able to translocate nucleic acid. This is

- 25 especially advantageous for gene therapy applications.

Thus according to one aspect of the present invention there is provided a conjugate comprising:

- (a) a first region comprising the homeodomain of antennapedia or a variant thereof; and
- (b) a second region not naturally associated with the first region; and wherein at least the first region is non-denatured.

5

According to one embodiment the first and second regions are associated via for example a disulfide bond, or the second region may comprise a nucleic acid binding domain, preferably further comprising nucleic acid. This embodiment may be seen as a protein/nucleic acid complex.

10

According to a further embodiment the conjugate is in the form of a fusion protein. In this embodiment, preferably the second region is a functional or regulatory protein, or an antigen.

15

- According to a further aspect of the present invention there is provided a conjugate comprising:
- (a) a first region comprising the homeodomain of antennapedia or a variant thereof operably linked to
- (b) a second region not naturally associated with the first region comprising a protein of at least 100 amino acids.

20

By "operably linked" we include that the first and second region are linked such that the second region is able to translocate a cell membrane.

25

Preferably the whole conjugate is non-denatured.

By non-denaturing we do not necessarily imply a specific non-denaturing step; although this may be the case.

Denaturation is the process by which the three-dimensional shape of the protein molecule is changed from its native state without rupture of peptide bonds. It can also include disulfide bond rupture or chemical modification of certain groups in the protein if these processes are also accompanied by changes in its overall three-dimensional structure.

We also do not exclude renatured proteins or nucleic acid in which the denatured protein or nucleic acid is returned to the conformation it maintained before denaturation. Reversible denaturation is generally brought about by disulfide 10 reducing agents, and urea, for nucleic acid by heat and salts.

For the avoidance of doubt, by native state we mean the form in which the protein or nucleic acid occurs in the intact cell, its three-dimensional structure depending on formation of the appropriate hydrogen bonds.

15 Preferably the first region is at the amino terminal end of the second region.

According to another aspect of the present invention there is provided nucleic acid encoding the conjugate of the present invention.

20 According to another aspect of the present invention there is provided an expression vector comprising the nucleic acid of the present invention operably linked to a promoter.

25 According to yet another aspect of the present invention there is provided a host cell transformed with the expression vector of the present invention.

The development of an appropriate procedure for obtaining the conjugate from bacterial lysate was important. With conventional methods we found that exposure

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to denaturing reagents dramatically affect the translocation property of the *Antp* homeodomain. We also found that small changes in pH and differences in osmolarity affected its translocation property.

5 We have now found a general method for preparing the conjugate of the present invention, and in particular an appropriate procedure for purifying the first domain and fusion proteins from the bacterial lysate. This method allows conjugates to be obtained which will translocate a protein of at least 100 amino acids across the cell membrane.

10

Thus according to another aspect of the present invention there is provided a method for preparing a conjugate comprising:

(i) culturing the host cell according to the present invention under conditions
15 which provide for the expression of the conjugate from the expression vector within the host cell; and
(ii) recovering the conjugate, which recovery comprises fusing an amino acid tail to the conjugate, which tail is capable of binding to at least one substrate and not to another substrate, and wherein the conjugate is caused to bind via the tail to at least
20 one substrate such that components of the host cell do not bind to this substrate; and the conjugate is contacted with the other substrate such that the conjugate is not bound and remaining components of the host cell are bound to the other substrate.

According to yet another aspect of the present invention there is provided a method
25 for preparing a conjugate comprising:

(i) culturing a host cell, transformed with an expression vector comprising nucleic acid, operably linked to a promoter, encoding (a) a first region comprising the homeodomain of antennapedia or a mutant thereof; and (b) a second region not

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naturally associated with the first region comprising an amino acid sequence of at least 100 amino acids in length, under conditions which provide for expression of the conjugate from the expression vector within the host cell; and

- (ii) recovering the conjugate, which method comprises comprises fusing an
5 amino acid tail to the conjugate, which tail is capable of binding to at least one substrate and not to another substrate, and wherein the conjugate is caused to bind via the tail to at least one substrate such that components of the host cell do not bind to this substrate; and the conjugate is contacted with the other substrate such that the conjugate is not bound and remaining components of the host cell are bound to the
10 other substrate.

According to a further aspect of the present invention there is provided a method of purifying a conjugate comprising fusing an amino acid tail to the conjugate, which tail is capable of binding to at least one substrate and not to another substrate, and

- 15 wherein the conjugate is caused to bind via the tail to at least one substrate such that impurities do not bind to this substrate; and the conjugate is contacted with the other substrate such that the conjugate is not bound and remaining impurities are bound to the other substrate.

- 20 Thus in broad terms the present invention can be seen as the use of a tail attached to the conjugate which allows both positive and negative purification steps.

Preferably the amino acid tail is fused to the carboxy terminal end of the conjugate.

- 25 Preferably the amino acid tail comprises HHHHHHG.

Preferably the substrate is a nickel column or an antibody with affinity for the amino acid tail.

According to a preferred embodiment the conjugate is consecutively contacted with two substrates with which it has affinity via the amino acid tail.

Thus preferably both the nickel column and antibody are used. They may be used in
5 any order.

When the second region is a DNA binding domain, a complex with nucleic acid may be formed by mixing the conjugate formed in accordance with the invention with the nucleic acid.
10

Further aspects of the present invention include a conjugate prepared by the method of the present invention; a pharmaceutical composition comprising the conjugate of the present invention and the use of the conjugate of the present invention in the preparation of a medicament for the treatment of a disease or infection.
15

In accordance with the invention, standard molecular biology techniques may be used which are within the level of skill in the art. Such techniques are fully described in the literature. See for example; Sambrook *et al* (1989) Molecular Cloning; a laboratory manual; Hames and Glover (1985 - 1997) DNA Cloning: a practical approach, Volumes I- IV (second edition); Methods for the engineering of immunoglobulin genes are given in McCafferty *et al* (1996) "Antibody Engineering: A Practical Approach".
20

Various preferred features and embodiments of the present invention will now be
25 described by way of non-limiting example, and with reference to the accompanying drawings in which:

Figure 1 shows the structure and sequence of the antennapedia homeodomain obtainable from *Drosophila*; and

Figure 2 further shows two mutants, designated pAntp 50H and pAntp 40P2.

First Region

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The first region of the conjugate of the invention may comprise a natural or synthetic homeodomain of antennapedia.

The homeodomain of the *Antp* gene obtainable from *Drosophila* is shown in Figure 1
10 and in Seq ID No. 1. Sequences homologous to this homeodomain have been isolated from other organisms, including vertebrates, mammals and humans, and these are included in the present invention. The homeodomain may be prepared using standard techniques such as cloning using the procedure described in Joliet et al (1991) *Antennapedia homeobox peptide regulates neural morphogenesis. Proc. Natl. acad. Sci. 88* 1864-1868. As previously indicated differences in the sequences of such multicellular organisms are generally conservative in nature. However, this may not necessarily be the case and other such sequences are included in the present invention, and for example where the sequence identity is about 50% or more, e.g. 60%, 70%, 80% or 90%, with the sequence obtainable from *Drosophila*. Sequence 15
20 identity may be determined using such commercially available programmes as GAP.

In addition synthetic variants may be used provided that they retain the ability to translocate the membrane. Synthetic variants will generally differ from the naturally-occurring proteins by substitution, particularly conservative substitution. By 25 conservative amino acid changes we mean replacing an amino acid from one of the amino acid groups, namely hydrophobic, polar, acidic or basic, with an amino acid from within the same group. An example of such a change is the replacement of valine by methionine and vice versa. Other examples of conservative substitutions may be seen by reference to the following table:

ALIPHATIC	Non-polar	G A P
	Polar - uncharged	I L V
		C S T M
	Polar - charged	N Q
AROMATIC		D E
OTHER		R K
		H F W Y
		N Q D E

Such variants may be prepared using standard recombinant DNA techniques such as site-directed mutagenesis. Where insertions are to be made, synthetic DNA encoding the insertion together with 5' and 3' flanking regions corresponding to the naturally-occurring sequence either side of the insertion site. The flanking regions will contain convenient restriction sites corresponding to sites in the naturally-occurring sequence so that the sequence may be cut with the appropriate enzyme(s) and the synthetic DNA ligated into the cut. The DNA is then expressed in accordance with the invention to make the encoded protein. These methods are only illustrative of the numerous standard techniques known in the art for manipulation of DNA sequences and other known techniques may also be used.

The ability of a naturally occurring or synthetic sequence to translocate the membrane may be tested by routine methods known in the art and illustrated in the accompanying examples.

Some variants of the homeodomain which retain the ability to translocate the membrane have been reported in the art and these are included in the scope of the present invention, together with any which become available.

10

For example, EP-B-0 485 578 to CNRS discloses homeopeptides comprising the helix 3 sequence of pAntp, and these are incorporated herein by reference.

WO97/12912 also to CNRS discloses the actual sequence of the helix 3 of pAntp,

- 5 and variants thereof. These also are incorporated herein by reference. In particular, the 3 helix is said to have the sequence:

Arg-Gln-Ile-Lys-Ile-Trp-Phe-Gln-Asn-Arg-Arg-Met-Lys-Trp-Lys-Lys

The variants are said to have the sequence:

X1-X2-X3-X4-X5-X6-X7-X8-X9-X10-X11-X12-X13-X14-X15-X16

10 or

X16-X15-X14-X13-X12-X11-X10-X9-X8-X7-X6-X5-X4-X3-X2-X1

wherein each X represents an α -amino acid, with X6 representing tryptophane; said peptide comprising between 6 and 10 hydrophobic amino acids.

- 15 Other variants are disclosed in for example, Gehring W (1987) Homeo Boxes in the Study of Development. *Science* 236 1245-1252 discloses a homeodomain of 62 amino acids, i.e. with glu at position 0 and lys at position 61. Bloch-Gallego E *et al* (1993) Antennapedia Homeobox Peptide Enhances Growth and Branching of Embryonic Chicken Motoneurons In Vitro. *The Journal of Cell Biology* 120(2) 485-
20 492 discloses a mutant called pAntp40P2 that was still able to translocate through the motoneuron membrane and to reach the nucleus. In this mutant the leucine and threonine residues in positions 40 and 41 were replaced by two proline residues. Le Roux *et al* (1993) Neurotropic activity of the Antennapedia homeodomain depends on its specific DNA-binding properties. *Proc. Natl. Acad. Sci.* 90 9120-9124
25 discloses two mutants pAntp 50A and pAntp 40P2 as shown in Figure 2 which retain the ability to translocate through the neuronal membrane. Schutze-Redelmeier M-P et al (1996) *supra* disclose that a 16 amino acid C-terminal (third helix) segment has been used to address oligonucleotides and oligopeptides to the cytoplasm and nuclei of cells in culture.

However, whilst not wishing to be bound by any theory it is believed that for delivering the proteins of the present invention an amino acid sequence of about 60 is preferred.

5

Cleavable linker region

Preferably, the first and second regions are linked by a cleavable linker region this may be any region suitable for this purpose. Preferably, the cleavable linker region is

- 10 a protease cleavable linker, although other linkers, cleavable for example by small molecules, may be used. These include Met-X sites, cleavable by cyanogen bromide, Asn-Gly, cleavable by hydroxylamine, Asp-Pro, cleavable by weak acid and Trp-X celavable by, *inter alia*, NBS-skatole. Protease cleavage sites are preferred due to the milder cleavage conditions necessary and are found in, for example, factor Xa,
- 15 thrombin and collagenase. Any of these may be used. The precise sequences are available in the art and the skilled person will have no difficulty in selecting a suitable cleavage site. By way of example, the protease cleavage region targeted by Factor Xa is I E G R. The protease cleavage region targeted by Enterokinase is D D D D K. The protease cleavage region targeted by Thrombin is L V P R G. Preferably 20 the cleavable linker region is one which is targetted by endocellular proteases.

Second Region

Fusion protein

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The second region of the conjugate according to the invention may comprise any protein sequence of interest (hereinafter POI) which is not naturally associated with the first region. Usually this will mean that the POI will be found in nature encoded by a gene different from the gene encoding the first region. The second region may

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be from the same species as the first region, but are present in the conjugate of the invention in a manner different from the natural situation, or from a different species.

The second region of the present invention is preferably at least 100 amino acids in length. The present invention is particularly useful for longer sequences, e.g. at least 150, 200, 300, 400 or 1000 amino acids in length. For the avoidance of doubt the term "protein" as used herein also encompasses polypeptides of the required length; although by the term "polypeptide" we generally mean sequences of from 2 to 100 amino acids in length, usually 2 up to 60.

10

Protein/nucleic acid complex

Within the present invention the nucleic acid comprises any nucleic acid of interest (hereinafter NOI), which may be e.g. therapeutically active or a reporter gene. The

15 NOI may be DNA or RNA. In one embodiment the NOI is an oligonucleotide.

When the present invention relates to a protein/nucleic acid complex, the complex preferably further comprises a nucleic acid (or DNA) binding domain as part of the second region. The nucleic acid binding domain serves to mediate the specific, high

20 affinity and non-covalent interaction of the protein component with the nucleic acid component.

The nucleic acid binding domain may be an RNA binding domain, or preferentially, a DNA binding domain, e.g. the DNA binding domain of a transcription factor,

25 particularly a yeast or human transcription factor. Preferred is A GAL4 derivable domain, mediating the selective binding of the protein of the invention to the DNA sequence CGGAGGACAGTCCTCCG (Cavey *et al* J Mol Biol 209:423, 1989). Most preferably the DNA binding domain consists of GAL4 amino acids 2 to 147. A DNA

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binding domain may bind to single-stranded or to a double stranded DNA on the second domain.

Another suitable domain would be a histone.

5 Applications for the conjugates of the present invention include (there may be overlap between these applications):

1. Antigen delivery system. An antigen is any agent that when introduced into an immunocompetent animal stimulates the production of a specific antibody or

10 antibodies that can combine with the antigen. However, the antigen may need to be combined with a carrier to be able to stimulate antibody production. This is where the present invention comes in as it acts as a carrier by transporting the antigen from one side of the cell membrane to the other such that it can stimulate antibody production. By way of example, bacterial and viral antigens translocated by *Antp* in 15 the cell cytoplasm may be processed and associated with MHC class I molecules. This antigen processing and presenting pathway is known to activate specific CD8 cytotoxic lymphocytes.

2. Gene therapy. Gene therapy may include any one or more of: the addition,

20 the replacement, the deletion, the supplementation, the manipulation etc. of one or more nucleotide sequences in, for example, one or more targeted sites - such as targeted cells. If the targeted sites are targeted cells, then the cells may be part of a tissue or an organ. General teachings on gene therapy may be found in Molecular Biology (Ed Robert Meyers, Pub VCH, such as pages 556-558).

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By way of further example, gene therapy can also provide a means by which any one or more of: a nucleotide sequence, such as a gene, can be applied to replace or supplement a defective gene; a pathogenic nucleotide sequence, such as a gene, or

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expression product thereof can be eliminated; a nucleotide sequence, such as a gene, or expression product thereof, can be added or introduced in order, for example, to create a more favourable phenotype; a nucleotide sequence, such as a gene, or expression product thereof can be added or introduced, for example, for

- 5 selection purposes (i.e. to select transformed cells and the like over non-transformed cells); cells can be manipulated at the molecular level to treat, cure or prevent disease conditions - such as cancer (Schmidt-Wolf and Schmidt-Wolf, 1994, Annals of Hematology 69:273-279) or other disease conditions, such as immune, cardiovascular, neurological, inflammatory or infectious disorders;
- 10 antigens can be manipulated and/or introduced to elicit an immune response, such as genetic vaccination. In a particularly preferred embodiment, *Antp* may be used to introduce functional proteins in the cytoplasm of genetically deficient cell types.

3. Cancer therapy. *Antp* may be used to transport into cancer cells molecules

- 15 that are transcription factors and are able to restore cell cycle control or induce differentiation. For example it is understood that many cancer cells would undergo apoptosis if a functional P-53 molecule is introduced into their cytoplasm. The present invention may be used to deliver such gene products.

- 20 4. Develop antibacterial and antiviral measures. For example, *Antp* may be used to transport in the cytoplasm of infected cells recombinant antibodies or DNA binding molecules and which interfere with a crucial step of bacterial and viral replication.

- 25 We have now generated recombinant chimeric proteins constituted by the homeodomain of antennapedia fused either to the 85A antigen of *Mycobacterium tuberculosis* or to a single chain recombinant antibody directed against the malaria antigen Pb21.

5. Use in expression systems. For example, it is desirable to express exogenous proteins in eukaryotic cells so that they get processed correctly. However, many exogenous proteins are toxic to eukaryotic cells. In manufacturing exogenous

5 proteins it is therefore desirable to achieve temporal expression of the exogenous protein. The system may therefore be used in connection with an inducible promoter for this or any other application involving such a system. In one such system, the loxP/cre system, the eukaryotic cell contains a gene encoding the protein of interest, functionally linked to a repressor. Thus, in the normal situation, the protein is not
10 expressed. The *Antp* homeodomain may be fused to Cre recombinase which will excise the repressor gene. Application of this conjugate to the eukaryotic cell will thus cause expression of the protein.

6. Receptor-mediated signalling. A receptor is a cellular macromolecule that

15 undergoes combination, with a hormone, neurotransmitter, drug or intracellular messenger to initiate a change in cell function. Receptors are concerned directly and specifically in chemical signalling between and within cells. By receptor-mediated we mean a response that requires the intermediary action of a receptor. Thus the present invention which is required to initiate a change in cell function.

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7. Protein sorting.

8. DNA synthesis.

25 The invention is particularly important because it transports to the nuclei of cells, functional transcription factors which interfere with the endogenous expression of genes and/or which interfere with the process of viral or bacterial replication.

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Suitable NOI coding sequences and POIs include those that are of therapeutic and/or diagnostic application such as, but are not limited to: sequences encoding or POIs which are cytokines, chemokines, hormones, antibodies, engineered immunoglobulin-like molecules, a single chain antibody, conjugates, enzymes,

- 5 immune co-stimulatory molecules, immunomodulatory molecules, anti-sense RNA, a transdominant negative mutant of a target protein, a toxin, suchas endotoxin A, Colicin A, d-endotoxin, diphtheria toxin, Bacillus anthrox toxin, Cholera toxin, Pertussis toxin, E. coli toxins, Shigatoxin or a Shiga-like toxin, a conditional toxin, an antigen, a tumour suppressor protein and growth factors, membrane 10 proteins, vasoactive proteins and peptides, anti-viral proteins and ribozymes, and derivatives therof (such as with an associated reporter group). When included, such coding sequences may be typically operatively linked to a suitable promoter, which may be a promoter driving expression of a ribozyme(s), or a different promoter or promoters.

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The conjugate of the present invention may be used to deliver one or more NOI(s) or POI(s) useful in the treatment of the disorders listed in WO-A-98/05635. For ease of reference, part of that list is now provided: cancer, inflammation or inflammatory disease, dermatological disorders, fever, cardiovascular effects,

- 20 haemorrhage, coagulation and acute phase response, cachexia, anorexia, acute infection, HIV infection, shock states, graft-versus-host reactions, autoimmune disease, reperfusion injury, meningitis, migraine and aspirin-dependent anti-thrombosis; tumour growth, invasion and spread, angiogenesis, metastases, malignant, ascites and malignant pleural effusion; cerebral ischaemia, ischaemic 25 heart disease, osteoarthritis, rheumatoid arthritis, osteoporosis, asthma, multiple sclerosis, neurodegeneration, Alzheimer's disease, atherosclerosis, stroke, vasculitis, Crohn's disease and ulcerative colitis; periodontitis, gingivitis; psoriasis, atopic dermatitis, chronic ulcers, epidermolysis bullosa; corneal

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ulceration, retinopathy and surgical wound healing; rhinitis, allergic conjunctivitis, eczema, anaphylaxis; restenosis, congestive heart failure, endometriosis, atherosclerosis or endosclerosis.

- 5 In addition, or in the alternative, the conjugate of the present invention may be used to deliver one or more NOI(s) or POI(s) useful in the treatment of disorders listed in WO-A-98/07859. For ease of reference, part of that list is now provided: cytokine and cell proliferation/differentiation activity; immunosuppressant or immunostimulant activity (e.g. for treating immune deficiency, including infection
- 10 with human immune deficiency virus; regulation of lymphocyte growth; treating cancer and many autoimmune diseases, and to prevent transplant rejection or induce tumour immunity); regulation of haematopoiesis, e.g. treatment of myeloid or lymphoid diseases; promoting growth of bone, cartilage, tendon, ligament and nerve tissue, e.g. for healing wounds, treatment of burns, ulcers and periodontal
- 15 disease and neurodegeneration; inhibition or activation of follicle-stimulating hormone (modulation of fertility); chemotactic/chemokinetic activity (e.g. for mobilising specific cell types to sites of injury or infection); haemostatic and thrombolytic activity (e.g. for treating haemophilia and stroke); antiinflammatory activity (for treating e.g. septic shock or Crohn's disease); as antimicrobials;
- 20 modulators of e.g. metabolism or behaviour; as analgesics; treating specific deficiency disorders; in treatment of e.g. psoriasis, in human or veterinary medicine.

In addition, or in the alternative, the conjugate of the present invention may be used to deliver one or more NOI(s) or POI(s) useful in the treatment of disorders listed in WO-A-98/09985. For ease of reference, part of that list is now provided: macrophage inhibitory and/or T cell inhibitory activity and thus, anti-inflammatory activity; anti-immune activity, i.e. inhibitory effects against a cellular and/or

humoral immune response, including a response not associated with inflammation; inhibit the ability of macrophages and T cells to adhere to extracellular matrix components and fibronectin, as well as up-regulated fas receptor expression in T cells; inhibit unwanted immune reaction and inflammation including arthritis,

- 5 including rheumatoid arthritis, inflammation associated with hypersensitivity, allergic reactions, asthma, systemic lupus erythematosus, collagen diseases and other autoimmune diseases, inflammation associated with atherosclerosis, arteriosclerosis, atherosclerotic heart disease, reperfusion injury, cardiac arrest, myocardial infarction, vascular inflammatory disorders, respiratory distress
- 10 syndrome or other cardiopulmonary diseases, inflammation associated with peptic ulcer, ulcerative colitis and other diseases of the gastrointestinal tract, hepatic fibrosis, liver cirrhosis or other hepatic diseases, thyroiditis or other glandular diseases, glomerulonephritis or other renal and urologic diseases, otitis or other oto-rhino-laryngological diseases, dermatitis or other dermal diseases, periodontal diseases or other dental diseases, orchitis or epididymo-orchitis, infertility, orchidal trauma or other immune-related testicular diseases, placental dysfunction, placental insufficiency, habitual abortion, eclampsia, pre-eclampsia and other immune and/or inflammatory-related gynaecological diseases, posterior uveitis, intermediate uveitis, anterior uveitis, conjunctivitis, chorioretinitis, uveoretinitis,
- 15 optic neuritis, intraocular inflammation, e.g. retinitis or cystoid macular oedema, sympathetic ophthalmia, scleritis, retinitis pigmentosa, immune and inflammatory components of degenerative fundus disease, inflammatory components of ocular trauma, ocular inflammation caused by infection, proliferative vitreo-retinopathies, acute ischaemic optic neuropathy, excessive scarring, e.g. following glaucoma
- 20 filtration operation, immune and/or inflammation reaction against ocular implants and other immune and inflammatory-related ophthalmic diseases, inflammation associated with autoimmune diseases or conditions or disorders where, both in the central nervous system (CNS) or in any other organ, immune and/or inflammation
- 25

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suppression would be beneficial, Parkinson's disease, complication and/or side effects from treatment of Parkinson's disease, AIDS-related dementia complex HIV-related encephalopathy, Devic's disease, Sydenham chorea, Alzheimer's disease and other degenerative diseases, conditions or disorders of the CNS.

- 5 inflammatory components of strokes, post-polio syndrome, immune and inflammatory components of psychiatric disorders, myelitis, encephalitis, subacute sclerosing pan-encephalitis, encephalomyelitis, acute neuropathy, subacute neuropathy, chronic neuropathy, Guillain-Barre syndrome, Sydenham chora, myasthenia gravis, pseudo-tumour cerebri, Down's Syndrome, Huntington's

10 disease, amyotrophic lateral sclerosis, inflammatory components of CNS compression or CNS trauma or infections of the CNS, inflammatory components of muscular atrophies and dystrophies, and immune and inflammatory related diseases, conditions or disorders of the central and peripheral nervous systems, post-traumatic inflammation, septic shock, infectious diseases, inflammatory

15 complications or side effects of surgery, bone marrow transplantation or other transplantation complications and/or side effects, inflammatory and/or immune complications and side effects of gene therapy, e.g. due to infection with a viral carrier, or inflammation associated with AIDS, to suppress or inhibit a humoral and/or cellular immune response, to treat or ameliorate monocyte or leukocyte

20 proliferative diseases, e.g. leukaemia, by reducing the amount of monocytes or lymphocytes, for the prevention and/or treatment of graft rejection in cases of transplantation of natural or artificial cells, tissue and organs such as cornea, bone marrow, organs, lenses, pacemakers, natural or artificial skin tissue.

25 The present invention also provides a pharmaceutical composition for treating an individual by gene therapy, wherein the composition comprises a therapeutically effective amount of the conjugate of the present invention. The pharmaceutical composition may be for human or animal usage. Typically, a physician will

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determine the actual dosage which will be most suitable for an individual subject and it will vary with the age, weight and response of the particular individual.

- The composition may optionally comprise a pharmaceutically acceptable carrier,
5 diluent, excipient or adjuvant. The choice of pharmaceutical carrier, excipient or diluent can be selected with regard to the intended route of administration and standard pharmaceutical practice. The pharmaceutical compositions may comprise as - or in addition to - the carrier, excipient or diluent any suitable binder(s), lubricant(s), suspending agent(s), coating agent(s), solubilising agent(s), and other
10 carrier agents that may aid or increase entry into the target site (such as for example a lipid delivery system).

- Where appropriate, the pharmaceutical compositions can be administered by any one or more of: inhalation, in the form of a suppository or pessary, topically in the
15 form of a lotion, solution, cream, ointment or dusting powder, by use of a skin patch, orally in the form of tablets containing excipients such as starch or lactose, or in capsules or ovules either alone or in admixture with excipients, or in the form of elixirs, solutions or suspensions containing flavouring or colouring agents, or they can be injected parenterally, for example intracavernosally, intravenously,
20 intramuscularly or subcutaneously. For parenteral administration, the compositions may be best used in the form of a sterile aqueous solution which may contain other substances, for example enough salts or monosaccharides to make the solution isotonic with blood. For buccal or sublingual administration the compositions may be administered in the form of tablets or lozenges which can be
25 formulated in a conventional manner.

The delivery of one or more therapeutic genes or proteins according to the invention may be used alone or in combination with other treatments or components of the

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treatment. Diseases which may be treated include, but are not limited to: cancer, neurological diseases, inherited diseases, heart disease, stroke, arthritis, viral infections and diseases of the immune system. Suitable therapeutic genes include those coding for tumour suppressor proteins, enzymes, pro-drug activating enzymes, immunomodulatory molecules, antibodies, engineered immunoglobulin-like molecules, conjugates, hormones, membrane proteins, vasoactive proteins or peptides, cytokines, chemokines, anti-viral proteins, antisense RNA and ribozymes.

The conjugate may also contain one or more cytokine-encoding NOIs or cytokines.

- 10 Suitable cytokines and growth factors include but are not limited to: ApoE, Apo-SAA, BDNF, Cardiotrophin-1, EGF, ENA-78, Eotaxin, Eotaxin-2, Exodus-2, FGF-acidic, FGF-basic, fibroblast growth factor-10 (Marshall 1998 *Nature Biotechnology* 16: 129).FLT3 ligand (Kimura et al. (1997), Fractalkine (CX3C), GDNF, G-CSF, GM-CSF, GF- β 1, insulin, IFN- γ , IGF-I, IGF-II, IL-1 α , IL-1 β , IL-2, IL-3, IL-4, IL-5,
- 15 IL-6, IL-7, IL-8 (72 a.a.), IL-8 (77 a.a.), IL-9, IL-10, IL-11, IL-12, IL-13, IL-15, IL-16, IL-17, IL-18 (IGIF), Inhibin α , Inhibin β , IP-10, keratinocyte growth factor-2 (KGF-2), KGF, Leptin, LIF, Lymphotactin, Mullerian inhibitory substance, monocyte colony inhibitory factor, monocyte attractant protein (Marshall 1998 *ibid*), M-CSF, MDC (67 a.a.), MDC (69 a.a.), MCP-1 (MCAF), MCP-2, MCP-3, MCP-4,
- 20 MDC (67 a.a.), MDC (69 a.a.), MIG, MIP-1 α , MIP-1 β , MIP-3 α , MIP-3 β , MIP-4, myeloid progenitor inhibitor factor-1 (MPIF-1), NAP-2, Neurturin, Nerve growth factor, β -NGF, NT-3, NT-4, Oncostatin M, PDGF-AA, PDGF-AB, PDGF-BB, PF-4, RANTES, SDF1 α , SDF1 β , SCF, SCGF, stem cell factor (SCF), TARC, TGF- α , TGF- β , TGF- β 2, TGF- β 3, tumour necrosis factor (TNF), TNF- α , TNF- β , TNIL-1,
- 25 TPO, VEGF, GCP-2, GRO/MGSA, GRO- β , GRO- γ , HCC1, 1-309,

The conjugate of the present invention may comprise further suitable domains. These will be known to those skilled in the art. For example an endoplasmic reticulum

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retention signal functions to affect the intracellular routing of the internalized conjugate or protein/nucleic acid complex of the present invention. A suitable endoplasmic retention signal may be a mammalian endoplasmic reticulum retention signal.

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Also present may be a translocation domain which serves to enhance nucleic acid or protein escape from the cellular vesicle system and thus to augment nucleic acid transfer by this route. This domain may serve to reduce or avoid lysosomal degradation after internalization of the protein/nucleic acid into the target cell. Suitable

10 translocation domains are derivable from toxins, particularly bacterial toxins, such as exotoxin A, Colicin A, d-endotoxin, diphtheria toxin, Bacillus anthrox toxin, Cholera toxin, Pertussis toxin, E. coli toxin toxins, Shigatoxin or Shiga-like toxin.

15 The first domain of the present invention may be modified to target cell sites other than the nucleus.

Additionally, or alternatively, also present may be a target cell-specific binding domain recognising a cell surface structure, such as a receptor protein or surface antigen on the target cell.

20

Nucleic acids.

The invention also provides nucleic acid encoding the conjugates of the invention. These may be constructed using standard recombinant DNA methodologies. The 25 nucleic acid may be RNA or DNA and is preferably DNA. Where it is RNA, manipulations may be performed via cDNA intermediates. Generally, a nucleic acid sequence encoding the first region will be prepared and suitable restriction sites provided at the 5' and/or 3' ends. Conveniently the sequence is manipulated in a standard laboratory vector, such as a plasmid vector based on pBR322 or pUC19 (see

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below). Reference may be made to Molecular Cloning by Sambrook *et al.* (Cold Spring Harbor, 1989) or similar standard reference books for exact details of the appropriate techniques.

- 5 Nucleic acid encoding the second region may likewise be provided in a similar vector system. Sources of nucleic acid may be ascertained by reference to published literature or databanks such as Genbank.

10 Nucleic acid encoding the desired first or second region may be obtained from academic or commercial sources where such sources are willing to provide the material or by synthesising or cloning the appropriate sequence where only the sequence data are available. Generally this may be done by reference to literature sources which describe the cloning of the gene in question.

15 Alternatively, where limited sequence data are available or where it is desired to express a nucleic acid homologous or otherwise related to a known nucleic acid, exemplary nucleic acids can be characterised as those nucleotide sequences which hybridise to the nucleic acid sequences known in the art.

20 Stringency of hybridisation refers to conditions under which polynucleic acids hybrids are stable. Such conditions are evident to those of ordinary skill in the field. As known to those of skill in the art, the stability of hybrids is reflected in the melting temperature (T_m) of the hybrid which decreases approximately 1 to 1.5°C with every 1% decrease in sequence homology. In general, the stability of a hybrid is 25 a function of sodium ion concentration and temperature. Typically, the hybridisation reaction is performed under conditions of higher stringency, followed by washes of varying stringency.

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As used herein, high stringency refers to conditions that permit hybridisation of only those nucleic acid sequences that form stable hybrids in 1 M Na⁺ at 65-68 °C. High stringency conditions can be provided, for example, by hybridisation in an aqueous solution containing 6x SSC, 5x Denhardt's, 1 % SDS (sodium dodecyl sulphate), 0.1

- 5 Na⁺ pyrophosphate and 0.1 mg/ml denatured salmon sperm DNA as non specific competitor. Following hybridisation, high stringency washing may be done in several steps, with a final wash (about 30 min) at the hybridisation temperature in 0.2 - 0.1x SSC, 0.1 % SDS.

- 10 Moderate stringency refers to conditions equivalent to hybridisation in the above described solution but at about 60-62°C. In that case the final wash is performed at the hybridisation temperature in 1x SSC, 0.1 % SDS.

- 15 Low stringency refers to conditions equivalent to hybridisation in the above described solution at about 50-52°C. In that case, the final wash is performed at the hybridisation temperature in 2x SSC, 0.1 % SDS.

- It is understood that these conditions may be adapted and duplicated using a variety of buffers, e.g. formamide-based buffers, and temperatures. Denhardt's solution and
20 SSC are well known to those of skill in the art as are other suitable hybridisation buffers (see, e.g. Sambrook, *et al.*, eds. (1989) Molecular Cloning: A Laboratory Manual, Cold Spring Harbor Laboratory Press, New York or Ausubel, *et al.*, eds. (1990) Current Protocols in Molecular Biology, John Wiley & Sons, Inc.). Optimal hybridisation conditions have to be determined empirically, as the length and the GC
25 content of the probe also play a role.

Expression vectors and host cells.

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The nucleic acid encoding a conjugate according to the invention, or constituent part(s) thereof, can be incorporated into vectors for further manipulation. As used herein, vector (or plasmid) refers to discrete elements that are used to introduce heterologous DNA into cells for either expression or replication thereof. Selection and use of such vehicles are well within the skill of the artisan. Many vectors are available, and selection of appropriate vector will depend on the intended use of the vector, i.e. whether it is to be used for DNA amplification or for DNA expression, the size of the DNA to be inserted into the vector, and the host cell to be transformed with the vector. Each vector contains various components depending on its function (amplification of DNA or expression of DNA) and the host cell for which it is compatible. The vector components generally include, but are not limited to, one or more of the following: an origin of replication, one or more marker genes, an enhancer element, a promoter, a transcription termination sequence and a signal sequence.

15

Production of Antibodies.

Conjugates according to the invention may be used directly as immunogens, without the use of further adjuvants, to generate antisera and monoclonal antibodies.

20

In accordance with yet another embodiment of the present invention, there are provided antibodies specifically recognising and binding the conjugates according to the invention. More preferably, however, the antibodies are specific for the second region of the conjugates, that is the polypeptide which is fused to the gene product of the invention in order to achieve expression thereof. Advantageously, the second region of the conjugate is recognised by the antibodies when in its natural context. Thus, where the second region is an isolated peptide or domain from a larger protein, that peptide or domain is recognised by the antibodies of the invention in the context of the whole of the larger protein.

The invention moreover provides a method for preparing an immunoglobulin, comprising the steps of:

- 5 a) immunising an animal with a conjugate according to the present invention :
and
b) recovering immunoglobulin specific for a region of the conjugate from the
serum of the animal.
- 10 The antibodies (or immunoglobulins) may be isolated in the form of a crude
preparation, i.e. an antiserum, by affinity chromatography against the conjugate.

Expression and purification of *Antp*-conjugates

- 15 Whilst not wishing to be bound by any theory we believe the ability of *Antp*-
conjugates to translocate across the cell surface membrane may be very much
dependent on the conformation of the recombinant proteins. We could not observe
any translocation of the polypeptide by using either bacteria cell extracts or purified
proteins exposed to small amounts of detergent (ionic and non-ionic) or denaturating
20 agents (urea or guanidinium). This conformation dependent property has represented
a serious problem for using *Antp*-conjugates in both *in vitro* and *in vivo* experiments
as all reports so far published use either synthetic peptides or bacterial cell extracts.
We have overcome this limitation by purifying the *Antp*-conjugate under native
conditions in accordance with the invention.

- 25 Example 1 - We have introduced at the carboxyl-terminal end of the recombinant
protein a tail of six histidine which confers to those proteins a high affinity for nickel
ions. Bacteria cells expressing the recombinant proteins were lysed by repeated
exposure to ultrasounds. Cell lysates were centrifuged and loaded on a nickel

column (Quiagen). The protein bound to the column were eluted step wise by changing the pH of the buffer. Under these conditions the *Antp*-conjugates could be selectively enriched. We have estimated by gel electrophoresis that the *Antp*-conjugates contributed up to 75% of the material eluted from the nickel column in

- 5 the pH range between 4.0 to 5.0. The *Antp* recombinant proteins were further purified by affinity chromatography using the antibody 4 D 11 directed against the histidine tail. The recombinant proteins were eluted from the column by changing the pH of the buffer. Our results indicate that fractions eluted at pH 2.8 contained the recombinant proteins nearly free of any bacterial contaminants. These protein
10 fraction were then loaded on a polymixine column to selectively subtract LPS and related bacterial contaminants. Gel electrophoresis analysis indicated that the eluate of the polymixine column contained the recombinant protein devoid of bacterial contaminants (99.5% purity). Purified *Antp*-conjugates were tested for their ability to translocate across the cell surface membrane of different cell types. Our results
15 indicate that the purified conjugates Antp-85A are able to translocate across a variety of cell lines (HeLa, Hep-G2, P815 EL4) and human monocytes. The ability of *Antp* to vehiculate large molecules across the cell membrane was not affected by the particular amino-acid composition of the conjugates. A different Antp-fusion molecule, generated by using the malaria antigen Thrombospondin related
20 anonymous protein (TRAP) of Plasmodium berghei of about 150 amino acids (Robson KJ et al Mol Biochem Parasitol 1997; 84(1): 1-12), was translocated across the cell membrane as efficiently as the 85A constructs.

The monoclonal antibody 4 D11

- 25 The monoclonal antibody (mAb) 4 D 11 was found by screening in ELISA hydridomas generated from a mouse that was immunised with a recombinant protein containing a six histidine tail at its amino-terminal end. The antibody is available from Imperial College of Science, Technology and Medicine, Sherfield Building Exhibition Rd, London SW7 2AZ, UK c/o ic Innovations Ltd, 47 Princes Gate,

London SW7 2AZ, UK. A molecular characterisation of the epitope showed that this mAb recognises the amino acid sequence HHHHHHGS both at the amino and at the carboxyl terminal end of recombinant proteins. The antibody has an IgG1 isotype and can be easily purified on protein A column. Our results indicate that 4 D 11

- 5 recognises the recombinant proteins containing the HHHHHHGS in ELISA immunoblot, immuno-fluorescence. In addition purified 4 D 11 coupled to beads (affi-gel or CNBR activated sepharose) can be used to purify recombinant proteins under native conditions.

10 TBC Vaccine

The present invention may be used to provide a recombinant vaccine that is able to induce a humoral as well as a cell mediated helper and cytotoxic immune response against *M. tuberculosis* antigens. In order to elicit a MCH class I restricted 15 cytotoxic immune response, the antigens employed in the vaccine should have access to the cytoplasmic compartment in the presenting cells of the immunised organism. The 85A protein is one of the member of the antigenic complex 85 ABC and represents the most abundant molecular species secreted *M. tuberculosis* in the culture media and during bacterial infection. Moreover, the protective immune 20 response induced natural infection is mainly directed against the antigen 85A, this protein elicits the production of antibodies and cytokines and stimulate the cytotoxic T cells. This would reproduce the processing and presentation pathways occurring during the natural infection of *M. tuberculosis* and the immunisation procedures employing BCG. Whilst not wishing to be bound by theory, it is believed that the 25 development of such a vaccine would require molecular vehicles for translocating bacterial proteins across the cell membrane into the cytoplasmic compartment.

Example 2 - To demonstrate the possibility of using *Antp* as delivery system we have carried out the following experimental activities: (i) cloning of the 85A full length

sequence in expression plasmids; (ii) development of synthetic genes encoding chimeric proteins in which the homeodomain of Ant has been inserted at the amino terminal end of the proteins 85A; (iii) expression in *E. coli* and purification of the recombinant proteins; (iv) development of a specific antiserum against 85A; and
5 (v) analysis of the ability of Antp to vehiculate the mycobacterial polypeptides 85A into the cytoplasm of HeLa cell in culture.

A. Cloning of 85A coding sequence in the expression plasmid pDS56/RBSII

The full length 85A gene was amplified in PCR experiments using as template DNA extracted *M. tuberculosis* strain H37Rv. As primers for the PCR reaction we have used oligonucleotides deduced from the amino acid sequence of 85A. To facilitate cloning in the expression vectors the primer were designed to contain at their 5' end the restriction sites *Bam* HI and *Sal* I. The amplified sequence was cloned in the expression plasmid pDS56/RBSII thereby generating the plasmids pDS/85A. The
10 amplified sequences 85A was sequenced to rule out the presence of mutations introduced during the amplification reaction. The expression unit of the plasmid pDS 56 RBSII is under the control of IPTG inducible promotor and yields conjugates containing a stretch of six histidines at their amino- or carboxyl-terminal end. The presence of the histidines confer to the proteins a high affinity for nickel ions. The
15 recombinant protein 85A was expressed in *E. coli* and purified by affinity chromatography on nickel column. The yields and quality of the proteins recovered at the end of purification process were assessed by analysing the samples eluted from the affinity column with SDS acrylamide gel electrophoresis.
20

25 B. Development of synthetic genes encoding the chimeric proteins Antp-85A

The sequence encoding the homeodomain of Antennapedia was amplified in PCR experiments using as template the DNA extracted from embryonic cells of *D. melanogaster*. The homeodomain encompasses the sequence of *Antp* from amino acid 297 to 356 and is encoded by a gene sequence that is not interrupted by introns.

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As primers we have employed oligonucleotides deduced on the basis of the homeodomain sequences and containing the Bam HI restriction site at their 5' end. The PCR product was sequenced to rule out the presence of mutations introduced by PCR in the sequence of the homeodomain. The restriction site Bam HI allowed the cloning of the PCR product in the correct reading frame in front of the sequence of 85A in the plasmid pDS/85A encoded the chimeric protein Antp-85A. This protein is characterised by the presence of the homeodomain of *Antp* and the histidine tail at its amino- and carboxyl-terminal ends respectively. The sequence of *Antp* was introduced at the amino-terminal end to facilitate the correct folding of this domain in the context of chimeric protein sequences. The expression of the chimeric protein Antp-85A was assessed by analysing the bacterial lysate on SDS acrylamide gels. In immunoblot experiments the serum s-85A recognised molecules migrating with the molecular weight predicted for Antp-85A.

15 C. Functional analysis of the chimeric proteins Antp-85A

To assess the ability of the chimeric protein to translocate across the cell membrane increasing amounts of purified Antp-85A were added to HeLa cells in culture. As control the cells were incubated with recombinant protein 85A that lacks the homeodomain of *Antp* at its amino-terminal end. After 3 hours of incubation at 37°C, the cells were fixed in formaldehyde and analysed with the antiserum s-85A to reveal the translocation of 85A epitopes into the cytoplasm. In immunofluorescence experiments the serum s-85A did not react with HeLa cells that were previously incubated with bacterial lysates containing 85A. This finding would indicate that these polypeptides are not able to cross the cell membrane. On the contrary the antiserum s-25 showed a clear cytoplasmic reactivity on HeLa cells that were incubated with the chimeric protein Antp-85A. This last result strongly indicated that the 85A epitopes were localised within the cytoplasm of cells incubated with Antp-85A thus indicating that the homeodomain of *Antp* has conferred to the chimeric proteins the ability to translocate across the superficial membrane. These

findings all together would indicate that the strategy employed to vehiculate antigens into the cytoplasm across the cell membrane can be used to develop an experimental vaccine against tuberculosis. It is anticipated that the purified chimeric proteins Antp-85A may elicit in the immunised animals both a humoral and cell mediated helper and cytotoxic immune response.

Example 3 - P/cre regulated expression of proteins in eukaryotic cells

The Cre recombinase is a phage derived enzyme that cuts double strand DNA at specific sites (LoxP). When two of these sites are present in opposite orientation the intervening DNA sequence is excised. The Cre recombinase has been shown to work in distantly related organisms such as bacteria, yeast and mammalian cells. *Antp* may be used to deliver the *Cre* recombinase inside the cell nucleus and for manipulating genomic DNA at precise locations in a temporal regulated manner.

One of the possible applications of *Antp-Cre* fusion is the development of highly regulated systems for expressing genes in eukaryotic cells. Transformation vectors can be designed for expressing genes under the transcriptional control of a promoter containing, within two LoxP sites, DNA elements functioning as target sequences for transcription repressor factors. In the absence of Cre recombinase the presence of the repressor sequence would not allow the promoter to transcribe the genes cloned in the vector. By adding the *Antp-Cre* conjugate to cells the repressor sequence may be excised from the promoter thus allowing transcription to start. This type of expression vector may be extremely useful for studying the function of eukaryotic genes as well as for expressing biologically active molecule in large amounts.

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Example 4 Control of viral expression by antennapedia delivery of recombinant transcription factors

We further investigate *Antp* delivery technology by assessing its ability to transport a functional form of the suppressive transcription factor Oct-2.4 into the nuclei of cultured cells. This factor is a member of the POU (Pit-Oct-Unc) family of transcription factors and originates from an alternative splicing of the Oct-2 gene transcript which only occurs in neuronal cells.

5 Oct-2.4 lacks the activation domain located at the C-terminal end of Oct-2 though it retains the ability to bind to the octamer DNA regulatory element AGTCAAT. In contrast to Oct-2, which acts as a positive regulatory factors on B lymphocytes promoters, Oct-2.4 can function as transcription repressor. Notably, Oct-2.4 was shown to suppress the transcription of the herpes simplex virus (HSV) immediate 10 early (IE) genes by binding to an octamer related regulatory element in the IE genes promoter. Moreover, the ectopic expression of Oct 2.4 in HSV permissive cells down regulated the IE gene expression and inhibited the viral lytic cycle thus 15 indicating that this factor may play an important function in determining the resistance of neuronal cells to HSV replication. Recombinant conjugates containing the homeodomain of *Antp* at the amino-terminal end of Oct-2.4 cross the cell surface membrane , reach the cell nuclei and down regulate the viral IE promoters.

20 This proposal has the objective to assess the ability of h*Antp* to transport Oct-2.4 across the surface membrane of living cells. To achieve this objective we: i) generate h*Antp*-Oct-2.4 fusion proteins; ii) assess the ability of the fusion proteins to bind to HSV IE promoters and to translocate across the cell membrane; iii) analyse the 25 transcriptional activity of IE promoter constructs in cells treated with h*Antp*-Oct-2.4 fusion proteins. The experiments prove the feasibility of manipulating the expression of cellular and viral genes by using a non invasive protein delivery system and may lead to the development of a novel class of antiviral agents.

Methods

Development of hAntp -Oct-2.4 constructs: Oct-2.4 is amplified in PCR experiments using as template cDNA generated from human B lymphocytes mRNA. These cells have been previously shown to express the full length Oct-2 transcription factor.

- 5 The Oct-2.4 PCR product is sequenced and used to generate constructs expressing recombinant proteins in *E. Coli*. The sequence encoding the homeodomain of *Antp* from amino acid 297 to 356 has been amplified from *Drosophila* DNA. This sequence has been modified by *in situ* mutagenesis to abolish its DNA binding properties and shown to translocate across the surface membrane of living cells. The
- 10 hAntp sequence is cloned at the 5' end of the Oct -2.4 gene thus generating the construct hAntp -Oct-2.4 that will contain hAntp at the amino-terminal end of Oct-2.4. hAntp -Oct-2.4 has a molecular weight of about 40 kDa which falls within the size range of proteins translocated by the homeodomain of *Antp*. The functional properties of Oct-2.4 will not be affected by the insertion of the hAntp sequence .
- 15 Structural analysis indicates that the suppressive and the DNA binding activity of Oct-2.4 are mediated by a 42 amino acid sequence (residues 141 -182) and the POU domain respectively which both retain their function in different structural contexts as well as in chimeric proteins. We also generate the construct hAntp-ΔOct-2.4 which has been designed to encode a deleted Oct-2.4 factor
- 20 encompassing the suppressive 42 amino acid sequence and the POU DNA binding domain. hAntp-ΔOct-2.4 will retain its suppressive transcriptional activity and translocate across the cell membrane more efficiently than hAntp-Oct-2.4 because of its reduced molecular size
- 25 **Expression of Oct-2.4 and hAntp-Oct- 2.4 chimeric proteins in *E. coli*:** hAntp-Oct-2.4, hAntp-ΔOct-2.4 and Oct-2.4 are expressed in *E. coli* using a vector that inserts a six histidine tail at their carboxyl-terminal ends. The presence of the His tail will allow the purification of the recombinant proteins by nickel chelate chromatography.

Experimental evidence indicates that proteins with this structural organisation are efficiently transported across the cell membrane.

To avoid the accumulation of incorrectly folded proteins we insert, at the amino-terminal end of h*Antp*-Oct-2.4, h*Antp*-ΔOct-2.4 and Oct-2.4 the signal sequence of

- 5 Pel B which directs the secretion of recombinant proteins into the bacteria periplasm. This strategy has proven to be very convenient to recover functional heterologous proteins in *E. coli*. The expression of recombinant proteins in *E. coli* are assessed in immunoblot experiments by using the MAb 4D11 which recognises the histidine tail.

- 10 Membrane translocation and DNA binding property analysis: The DNA binding properties of the recombinant proteins h*Antp*-Oct-2.4, h*Antp*-ΔOct-2.4 and Oct-2.4 are assessed in gel retardation assays using oligonucleotides encompassing the natural and the related HSV octamer sequence. The specificity of the interaction will
15 be investigated in competition assays using wild type and mutated octamer oligonucleotides. The h*Antp*-Oct-2.4 constructs are further investigated for their ability to cross the surface membrane of several cell lines including the BHK cells (ATCC: CRL-1632) which are susceptible to HSV infection and lack Oct-2 transcription factors. BHK cells are incubated with affinity purified h*Antp*-Oct-2.4,
20 h*Antp*-ΔOct-2. Control experiments are carried out by incubating the cells with recombinant Oct-2.4 and unrelated h*Antp*-fusion proteins. The presence and the distribution of the recombinant constructs in the treated cells are analysed in immunofluorescence experiments. BHK cells incubated with h*Antp*-Oct-2.4, h*Antp*-ΔOct-2.4 and Oct-2.4 are fixed, permeabilised and processed for
25 immunofluorescence using MAb 4 D11.

Analysis of the transcriptional activity of h*Antp*-Oct-2.4 and h*Antp*-ΔOct-2.4: The suppressive transcriptional activity of h*Antp*-Oct-2.4 and h*Antp*-ΔOct-2 are assessed on BHK cells transformed with a HSV IE promoter reporter construct. A DNA

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sequence encompassing the IE gene 3 promoter from nucleotide -330 to +33 will be linked to the luciferase (*Luc*) gene in the transformation construct IE3-Luc. The selected DNA sequences contains the octamer related regulatory sequence TAATGARAT and is transcriptionally active in BHK cells. The IE3-Luc construct 5 also contains a G418 resistance gene (*neo*) transcription unit consisting of: i) the MMTV-LTR retroviral promoter-ii) the *neo* gene- and iii) the SV40 splicing and polyadenylation sites. Transformed BHK cell clones are selected with genetin(G418) and analysed by southern blot. Experimental evidence indicates that the HSV IE3 promoter will constitutively express high level of luciferase in IE3-Luc 10 transformed BHK cells. To assess the ability of h*Antp*-Oct-2.4, h*Antp*-ΔOct-2.4 to suppress the transcriptional activity of the HSV IE3 promoter, IE3-Luc transformed BHK cells are cultured either in the absence or in the presence of different concentration of affinity purified h*Antp* chimeric proteins. As control the transformed cells are incubated with recombinant Oct-2.4 and unrelated h*Antp* 15 fusion proteins. The transcriptional level of the HSV IE3 promoter are assessed by measuring the luciferase activity in BHK cell extracts collected at different time points after treatment. Functional h*Antp*-Oct-2.4, h*Antp*-ΔOct-2.4 chimeric protein translocated into the cell nucleus of IE3-Luc transformed BHK will significantly down regulate the baseline activity of the HSV IE3 promoter.

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The relevance of the proposed experiments goes beyond the development a delivery system potentially able to interfere with the lytic cycle of HSV. The homeodomain of *Antp* has the ability to transport across the cell surface membrane a range of functional proteins of great biological and medical relevance.

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CLAIMS

1. A conjugate comprising:
 - (a) a first region comprising the homeodomain of antennapedia or a functional variant thereof; and
 - 5 (b) a second region not naturally associated with the first region; and wherein at least the first region is non-denatured.
2. A conjugate according to claim 1, wherein the first and second regions are associated via a disulfide bond.
- 10 3. A conjugate according to claim 1, in the form of a fusion protein.
4. A conjugate according to any preceding claim, wherein the second region comprises a protein of at least 100 amino acids.
- 15 5. A conjugate according to claim 4, wherein the second region is a functional or regulatory protein.
6. A conjugate according to claim 4, wherein the second region is an antigen.
7. A conjugate according to claim 4, wherein the second 20 region is a DNA binding domain.
8. A conjugate according to claim 7, wherein the second region further comprises an NOI.
9. A conjugate according to claim 1 or claim 2, wherein the second region comprises an NOI.
- 25 10. Nucleic acid encoding a conjugate according to any preceding claim.
11. An expression vector, comprising the nucleic acid of claim 10, operably linked to a promoter.
12. A host cell transformed with the expression vector of 30 claim 11.
13. A method for preparing a conjugate according to any of claims 1 to 9, comprising:
 - (i) culturing a host cell according to claim 12 under conditions which provide for the expression of the conjugate within the host cell; and
 - 35 (ii) recovering the conjugate by affinity purification under non-denaturing conditions.

14. A method according to claim 13, wherein the conjugate comprises an amino acid tail that binds to an immobilised substrate.
- 5 15. A conjugate prepared according to the method of claim 13 or claim 14.
16. A pharmaceutical composition comprising the conjugate of any one of claims 1 to 9 or claim 15, in combination with a pharmaceutically-acceptable carrier.
- 10 17. A pharmaceutical composition according to claim 16, in the form of a vaccine.
18. Use of the conjugate of any one of claims 1 to 9 or claim 15, in the preparation of a medicament for the treatment or prevention of cancer, a genetic disease and bacterial or viral infection.
- 15 19. A conjugate according to any one of claims 1 to 9 or claim 15, for use in an expression system.

Abstract of the Disclosure

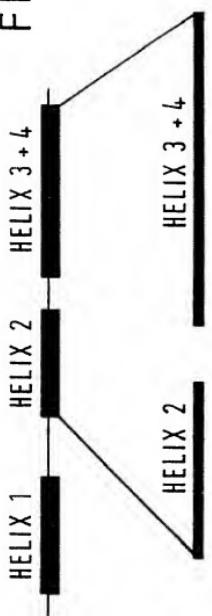
The subject invention pertains to a conjugate comprising: (a) a first region comprising the homeodomain of antennapedia or a variant thereof; and (b) a second region not naturally associated with the first region. In one embodiment, the second region of the conjugate comprises a protein of at least 100 amino acids.

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FIG. 1.



LIRRRIEIAHALCLTERQIKIWFKQNRMMWKKKEN pAntp

FIG. 2.

LIRRRIEIAHALCLTERQIKIWFKQNRMMWKKKEN pAntp
Y-----A-----pAntp 50A
P-----P-----pAntp 40P2

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As a below named inventor, I hereby declare that my residence, post office address and citizenship are as stated below next to my name; I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of subject matter which is claimed and for which a patent is sought on an invention entitled
CONJUGATES THAT CONTAIN THE HOMEODOMAIN OF ANTENNA PEDIA

the specification of which is attached hereto or

was filed on 02 SEP 1998 as United States Application Number or PCT International Application Number PCT/GB98/02628 and was amended on 24 SEP 1999 (if applicable)

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above. I acknowledge the duty to disclose information which is material to patentability as defined in 37 CFR 1.56. I hereby claim foreign priority benefits under 35 U.S.C. 119(a)-(d) or 365(b) of any foreign application(s) for patent or inventor's certificate, or 365(a) of any PCT international application which designated at least one country other than the United States of America, listed below and have also identified below, by checking the box, any foreign application for a patent or inventor's certificate, or PCT international application having a filing date before that of the application on which priority is claimed:

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			<input type="checkbox"/>	<input type="checkbox"/>	NO
9718609.2	GB	02 SEP 1997	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
			<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
			<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

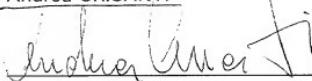
As a named inventor, I hereby appoint the following registered practitioner(s) to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith:
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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under 18 U.S.C 1001 and that such willful false statements may jeopardise the validity of the application or any patent issued thereon.

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